

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
14 February 2002 (14.02.2002)

PCT

(10) International Publication Number
WO 02/12153 A1

(51) International Patent Classification⁷: **C07C 17/395**,
17/383, 17/386, 17/389, 19/08

(21) International Application Number: PCT/EP01/10064

(22) International Filing Date: 10 August 2001 (10.08.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:

| | | |
|----------|-----------------------------|----|
| 00/10549 | 10 August 2000 (10.08.2000) | FR |
| 00/10548 | 10 August 2000 (10.08.2000) | FR |
| 00/10547 | 10 August 2000 (10.08.2000) | FR |
| 00/10546 | 10 August 2000 (10.08.2000) | FR |
| 01/03193 | 7 March 2001 (07.03.2001) | FR |
| 01/08910 | 4 July 2001 (04.07.2001) | FR |

(71) Applicant (for all designated States except US): **SOLVAY (SOCIÉTÉ ANONYME)** [BE/BE]; Rue du Prince Albert 33, B-1050 Bruxelles (BE).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **WILMET, Vincent** [BE/BE]; Rue Cour Boisacq 81, B-1301 Wavre (BE). **JANSSENS, Francine** [BE/BE]; Steenweg op Koningslo 106, B-1800 Vilvoorde (BE). **CASAUBON SEIGNOUR, Lionel** [FR/FR]; Rue Gay-Lussac 8, F-39500 Tavaux (FR). **KRAFFT, Philippe** [FR/FR]; Rue Gay-Lussac 11, F-39500 Tavaux (FR). **LAMBERT, Alain** [BE/BE]; Rue du Broux 41, B-1320 Beauvechain (BE). **BUYLE, Olivier** [BE/BE]; Rue des Chanterelles 86, B-6030 Goutroux (BE).

(74) Agents: **JACQUES, Philippe** et al.; SOLVAY (Société Anonyme), Département de la Propriété Intellectuelle, Rue de Ransbeek 310, B-1120 Bruxelles (BE).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Declarations under Rule 4.17:

- of inventorship (Rule 4.17(iv)) for US only
- of inventorship (Rule 4.17(iv)) for US only
- of inventorship (Rule 4.17(iv)) for US only
- of inventorship (Rule 4.17(iv)) for US only
- of inventorship (Rule 4.17(iv)) for US only

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: PROCESS FOR OBTAINING A PURIFIED HYDROFLUOROALKANE

(57) Abstract: Process for obtaining a hydrofluoroalkane which is purified of organic impurities, according to which the hydrofluoroalkane containing organic impurities is subjected to at least one purification treatment chosen from: (a) a treatment with chlorine in the presence of a initiator, (b) a reaction with hydrogen fluoride, (c) a distillation in which the purified hydrofluoroalkane is removed from the top of the distillation column or from the side, (d) an extractive distillation, (e) an adsorption onto a solid adsorbent, (f) a reaction with a compound containing oxygen, and (g) a reaction with a compound containing oxygen and a gas-phase reaction with a reagent capable of reacting with at least some of the organic impurities, with the exception of a reaction with elemental chlorine.



WO 02/12153 A1

Process for obtaining a purified hydrofluoroalkane

The present invention relates to a process for obtaining a purified hydrofluoroalkane chosen in particular from 1,1,1,3,3-pentafluoropropane and 1,1,1,3,3-pentafluorobutane.

5 Hydrofluoroalkanes such as 1,1,1,3,3-Pentafluoropropane and 1,1,1,3,3-pentafluorobutane may be used as replacements for (hydro)chlorofluoroalkanes, for example as blowing agents, as refrigerants or as solvents.

These hydrofluoroalkanes are typically manufactured by reacting a chloro or chlorofluoro precursor with hydrogen fluoride. The crude hydrofluoroalkanes
10 obtained in this reaction often contain impurities such as unconverted reagents, hydrogen chloride and olefinic impurities, in particular chlorofluoro olefins containing 3 or 4 carbon atoms.

Patent application WO-A-00/14040 describes a process for purifying 1,1,1,3,3-pentafluorobutane. According to this known process, it is possible to
15 reduce the fluorotrichloroethylene content in 1,1,1,3,3-pentafluorobutane by ionic chlorination in the presence of FeCl_3 , by hydrogenation in the presence of Pd/Rh on active charcoal or, in particular, by reaction with fluorine.

Patent application WO-A-97/37955 describes a process for purifying 1,1,1,3,3-pentafluoropropane of 1-chloro-3,3,3-trifluoropropene, in which a
20 photochlorination initiated with UV light of wavelength from 300 to 400 nm is carried out.

The Applicant has now found that hydrofluoroalkenes – that is to say olefins consisting solely of carbon, hydrogen and fluorine – are particularly difficult to remove when they are present as impurity in a hydrofluoroalkane, in
25 particular those comprising from 3 to 5 carbon atoms, most particularly those corresponding to the empirical formula $\text{C}_4\text{H}_4\text{F}_4$, present as impurity in particular in 1,1,1,3,3-pentafluorobutane. On account of the very low chemical reactivity observed, the removal by means of a chemical treatment of the hydrofluoroalkenes in hydrofluoroalkanes is liable to require prolonged treatment
30 times that are undesirable in an industrial process for manufacturing hydrofluoroalkane. In an extreme case, it would not be possible to go below a certain hydrofluoroalkene content.

None of the prior art documents addresses this specific problem.

It was consequently desirable to have available a process for purifying
35 hydrofluoroalkane, in particular 1,1,1,3,3-pentafluoropropane or 1,1,1,3,3-

pentafluorobutane, which allows an effective reduction of the content of olefinic impurities and in particular of hydrofluoroalkenes while at the same time using technical means that are simple to implement.

- The invention consequently relates to a process for obtaining a
- 5 hydrofluoroalkane comprising at least three carbon atoms, which is purified of organic impurities, according to which the hydrofluoroalkane containing organic impurities including (chloro)fluoro olefins is subjected to at least one purification treatment chosen from
- (a) a treatment with chlorine in the presence of an initiator
 - 10 (b) a reaction with hydrogen fluoride
 - (c) a distillation in which the purified hydrofluoroalkane is removed from the top of the distillation column or from the side
 - (d) an extractive distillation
 - (e) an adsorption onto a solid adsorbent
 - 15 (f) a reaction with a compound containing oxygen, and
 - (g) a gas-phase reaction with a reagent capable of reacting with at least some of the organic impurities, with the exception of a reaction with elemental chlorine.

The process according to the invention applies in particular to hydrofluoroalkanes such as 1,1,1,3,3,3-hexafluoropropane (HFC-236fa),

20 1,1,1,2,3,3,3-heptafluoropropane (HFC-227ea) and 1,1,1,2,3,4,4,5,5,5-decafluoropentane (HFC-43-10mee). Preferably, the hydrofluoroalkane comprises 3 or 4 carbon atoms. The process has been found to be particularly effective for purifying 1,1,1,3,3-pentafluoropropane or 1,1,1,3,3-pentafluorobutane, and most particularly the latter.

25 It has been found surprisingly that the process according to the invention allows an effective reduction of the content of organic impurities in the hydrofluoroalkane. In particular, 1,1,1,3,3-pentafluorobutane and 1,1,1,3,3-pentafluoropropane have physical and chemical stability under the conditions of the process according to the invention. The process according to the invention

30 may be carried out easily.

The organic impurities whose content may be reduced by means of the process according to the invention generally comprise 3 or 4 carbon atoms. They may also comprise at least one chlorine atom, such as chlorodifluoropropanes and chlorofluorobutanes or -butenes. They are in particular (chloro)fluoro olefins

35 containing 3 or 4 carbon atoms, such as monochlorotrifluorobutene isomers.

The process according to the invention is particularly suitable for effectively removing hydrofluoroalkenes as described above. The process according to the invention, which may be carried out easily, allows an effective reduction of the content of olefinic impurities present in 1,1,1,3,3-
5 pentafluoropropane or 1,1,1,3,3-pentafluorobutane.

In a first aspect of the process according to the invention, the hydrofluoroalkane containing olefinic impurities, in particular hydrofluoroalkenes comprising from 3 to 5 carbon atoms, is subjected to a treatment with chlorine in the presence of an initiator.

10 The treatment with chlorine serves to chlorinate the olefinic impurities in the hydrofluoroalkane containing olefinic impurities. These are notably (chloro)fluoro olefins containing 3 or 4 carbon atoms or, in particular, the hydrofluoroalkenes mentioned above.

The initiator serves to decompose the chlorine molecules by cleavage.
15 In a first variant of the first aspect of the process according to the invention, the the initiator is a free-radical initiator selected from an organic or inorganic initiator compound.

To promote the mixing of the hydrofluoroalkane containing olefinic impurities with the initiator compound, the first variant of the first aspect of the process according to the invention is preferably carried out in the liquid phase.
20

According to the invention, the free-radical initiator is often an organic compound. Among the organic compounds that are usually used are peroxide or diazo compounds. Peroxide compounds are used in particular. Among these, the ones chosen more particularly are diacyl peroxides, peroxydicarbonates, alkyl
25 peresters, peracetals, ketone peroxides, alkyl hydroperoxides and dialkyl peroxides. Diacyl peroxides or peroxydicarbonates are preferably selected. Excellent results have been obtained with dilauroyl peroxide, dibenzoyl peroxide or dicetyl peroxydicarbonate.

The free-radical initiator may also be an inorganic compound. In this case,
30 it is often chosen from hydrogen peroxide, percarbonates such as, in particular, sodium percarbonate, and perborates such as sodium perborate.

The initiator compound is preferably selected from compounds with a half-life from 0.1 to 3 hours and usually of about 1 hour at the temperature of the treatment with chlorine.

35 The initiator compound is generally used in a proportion of at least about 10 ppm by weight relative to the hydrofluoroalkane containing olefinic impurities.

At least about 20 ppm by weight of initiator compound are used. Even more particularly at least about 30 ppm by weight in particular. Most frequently, not more than about 10 000 ppm by weight of initiator compound are used relative to the hydrofluoroalkane containing olefinic impurities. Preferably, the amount of organic initiator compound does not exceed about 1 000 ppm by weight and even more preferably it does not exceed about 300 ppm by weight.

In the first variant of the first aspect of the process according to the invention, the chlorine may be used in the gas phase or in the liquid phase. It is introduced in excess amounts relative to all of the olefinic impurities to be chlorinated in the hydrofluoroalkane containing olefinic impurities. Generally, the chlorine is used in a proportion of more than 3 mol per mole of olefinic impurities and preferably at least about 4 mol per mole of olefinic impurities. Generally, it is not desirable to exceed about 40 mol of chlorine per mole of olefinic impurities. It is preferable to limit the amount used so that virtually all of the chlorine can react and is not found in unchanged form after the present purification treatment. Preferably, the amount does not exceed about 15 mol per mole of olefinic impurities and even more preferably this ratio does not exceed about 12.

In the first variant of the first aspect of the process according to the invention, the treatment with chlorine may be carried out over a wide temperature range. In particular, the treatment with chlorine is carried out at a temperature of at least about 40°C and even more particularly of at least about 60°C. Higher temperatures allow a faster conversion of the unsaturated compounds. However, this results in a correlative increase in pressure, of which account needs to be taken. Preferably, the treatment temperature does not exceed about 150°C and even more preferably it does not exceed about 100°C. Excellent results have been obtained when the treatment with chlorine is carried out in the regions of 60 to 100°C.

In the first variant of the first aspect of the process according to the invention, the treatment with chlorine may be carried out at the autogenous pressure or a higher pressure generated, for example, by introducing an inert gas. In general, the treatment is carried out at a pressure which does not exceed about 5 MPa and preferably 2 MPa. Pressures from about 0.2 to about 1.0 MPa are suitable for use.

These correlated conditions of high temperature and high pressure which are allowed for the treatment with chlorine contribute towards the fast and effective removal of the olefinic impurities. In the first variant of the first aspect

of the process according to the invention, the duration of the treatment with chlorine may be from about 1 to about 120 minutes. Preferably, the duration of the treatment with chlorine is not more than about 60 minutes.

According to an advantageous embodiment of the first variant of the first aspect of the process according to the invention, the initiator compound is introduced into the hydrofluoroalkane containing olefinic impurities before the chlorine. In a preferred implementation variant of this embodiment of the invention, the chlorine is introduced into the hydrofluoroalkane at a temperature close to the treatment temperature. In a particularly preferred implementation variant of this embodiment of the invention, the initiator compound is also introduced into the hydrofluoroalkane at a temperature close to the treatment temperature.

In a second variant of the first aspect of the process according to the invention, the initiator, which is a free-radical initiator, is an electromagnetic radiation comprising at least one fraction of wavelengths less than 280 nm.

It has been found, surprisingly, that this variant of the first aspect of the process according to the invention is particularly effective for reducing to an acceptable level the amount of the hydrofluoroalkenes which may be contained in a hydrofluoroalkane, quickly and without substantial degradation of the hydrofluoroalkane. This variant of the first aspect of the process according to the invention allows the chlorine to be used in the presence of a broader spectrum of wavelengths than in the known processes. This variant of the first aspect of the process according to the invention allows an accelerated purification operation, an efficient destruction of the chlorine-free impurities and an improved use of energy.

This variant of the first aspect of the process according to the invention is particularly suitable for purifying 1,1,1,3,3-pentafluorobutane of hydrofluoroalkenes. It is more particularly suitable for purifying 1,1,1,3,3-pentafluorobutane of hydrofluoroalkenes of empirical formula $C_4H_4F_4$, in particular *E*-CF₃-CH=CF-CH₃, *Z*-CF₃-CH=CF-CH₃, *E*-CF₃-CH=CH-CH₂F, *Z*-CF₃-CH=CH-CH₂F, *E*-CF₃-CH₂-CH=CHF, *Z*-CH₃-CH₂-CH=CHF and/or CF₃-CH₂-CF=CH₂. The process is particularly suitable for purifying 1,1,1,3,3-pentafluorobutane of one or more hydrofluoroalkenes chosen from *E*-CF₃-CH=CF-CH₃, *Z*-CF₃-CH=CF-CH₃ and CF₃-CH₂-CF=CH₂.

It is understood that the first variant of the first aspect of the process according to the invention and any of the methods described below may also be

used to purify the hydrofluoroalkane, in particular 1,1,1,3,3-pentafluorobutane, of hydrofluoroalkenes as mentioned above. However, the second variant of the first aspect of the process according to the invention is, in the light of the advantages described above, most particularly preferred for purifying hydrofluoroalkanes of
5 hydrofluoroalkenes.

In the second variant of the first aspect of the process according to the invention, the energy of the fraction of wavelengths less than 280 nm is generally at least 5% of the total energy of the electromagnetic radiation. Preferably, the energy of the fraction of wavelengths less than 280 nm is at least 10% of the total
10 energy of the electromagnetic radiation. An electromagnetic radiation in which 100% of the energy is contained in the fraction of wavelengths less than 280 nm may even be used.

The wavelengths in the fraction of wavelengths less than 280 nm are preferably essentially at least 170 nm. In a more particularly preferred manner, the
15 wavelengths are essentially at least 180 nm. In one variant, the fraction of wavelengths comprises radiation of wavelength less than or equal to 270 nm, in particular 260 nm.

Sources of electromagnetic radiation comprising a fraction of wavelengths less than 280 nm are, for example, ultraviolet (UV) burners such as, in particular,
20 medium pressure or, preferably, low pressure, optionally doped, high power mercury vapour burners. Such burners are commercially available, for example from companies Hereaus or ABB. A monochromatic burner of suitable wavelength may also be used.

The separation between the burner and the reaction medium in which the
25 purification reaction is carried out is generally achieved with a translucent material which allows electromagnetic radiation of wavelength less than 280 nm to pass through. An example of such a material is quartz.

The intensity of the electromagnetic radiation is generally at least 0.01 W.h per kg of hydrofluoroalkane containing impurities, preferably at least 0.02
30 W.h.kg⁻¹ or even at least 0.05 W.h.kg⁻¹. The intensity of the electromagnetic radiation is generally not more than 5 W.h per kg of hydrofluoroalkane containing impurities and preferably not more than 3 W.h.kg⁻¹ or even not more than 2 W.h.kg⁻¹.

In the second variant of the first aspect of the process according to the
35 invention, the chlorine may be used in the gas phase or in the liquid phase. It is preferably used in the liquid phase.

The second variant of the first aspect of the process according to the invention may be carried out, for example, in a falling film photoreactor or in an immersed burner photoreactor.

In a first embodiment of the second variant of the first aspect of the process according to the invention, the chlorine is introduced in stoichiometric or excess amounts relative to the entirety of the olefinic impurities to be chlorinated in the hydrofluoroalkane containing impurities. In this embodiment, the chlorine is used in an amount of greater than or equal to about 1 mol per mole of olefinic impurities. The amount of chlorine is, in this embodiment, generally less than or equal to about 10 mol of chlorine per mole of olefinic impurities. Preferably, the amount does not exceed about 5 mol of chlorine per mole of olefinic impurities and even more preferably this ratio does not exceed about 2.

In a second embodiment of the second variant of the first aspect of the process according to the invention, the chlorine is introduced in amounts less than the entirety of the olefinic impurities to be chlorinated in the hydrofluoroalkane containing impurities. In this variant, the chlorine is preferably used in an amount of less than about 0.9 mol per mole of olefinic impurities. The amount of chlorine is, in this embodiment, generally greater than or equal to about 0.01 mol of chlorine per mole of olefinic impurities. Preferably, this amount is greater than or equal to about 0.1 mol of chlorine per mole of olefinic impurities. An amount of greater than or equal to about 0.5 mol of chlorine per mole of olefinic impurities is most particularly preferred.

In the second variant of the first aspect of the process according to the invention, the treatment with chlorine is generally carried out at a temperature of greater than or equal to -20°C . The temperature is often greater than or equal to 0°C . Preferably, the temperature is greater than or equal to about 10°C . In this variant, the treatment with chlorine is generally carried out at a temperature of less than or equal to 150°C . The temperature is often less than or equal to 100°C . Preferably, the temperature is less than or equal to about 60°C .

In the second variant of the first aspect of the process according to the invention, the pressure at which the treatment with chlorine is carried out is generally greater than or equal to about 1 bar. The pressure at which the treatment with chlorine is carried out is generally less than or equal to about 10 bar. Preferably, the pressure is less than or equal to about 5 bar.

In the second variant of the first aspect of the process according to the invention, the duration of the treatment with chlorine is generally greater than or

equal to 5 min. The duration of the treatment with chlorine is often greater than or equal to 10 min. Preferably, the duration of the treatment with chlorine is greater than or equal to 15 min. In the second variant of the first aspect of the process according to the invention, the duration of the treatment with chlorine is generally
5 less than or equal to 10 h. The duration of the treatment with chlorine is often less than or equal to 5 h. Preferably, the duration of the treatment with chlorine is less than or equal to about 3 h. In a particularly preferred manner, it does not exceed 2 h.

In a third variant of the first aspect of the invention the initiator is a low
10 amount of metal ion. The third variant is carried out preferably in the substantial absence of free-radical initiators. In particular it is preferably carried out in the substantial absence of electromagnetic radiation having a wavelength from 170 nm to 400 nm (UV-light). According to this variant, efficient elimination of hydrofluoroalkenes such as mentioned above may be achieved, without
15 substantial degradation of the desired hydrofluoroalkane. In this variant, no specific separation operation is required to separate the initiator from the hydrofluoroalkane. Alternatively, the initiator is separated easily by an optional distillation

The metal ion is preferably a lewis acid. It is preferably selected from ions
20 of group IIIa, IVa and b, Va and b, VIb and VIII metals of the Periodic Table of Elements (IUPAC 1970). In a particularly suitable manner, it is selected from ions of iron, nickel, aluminium, boron, titanium, chromium, zirconium, tantalum, tin or antimony. Iron ions are particularly preferred.

The amount of metal ion present in the treatment with chlorine is generally
25 at most 1000 ppm by weight relative to the hydrofluoroalkane containing organic impurities. The amount of metal ion is more frequently at most 100 ppm. The amount is preferably at most 50 ppm. An amount of metal ions of at most 30 ppm is particularly preferred. The amount of metal ion present in the treatment with chlorine is generally at least 0,01 ppm by weight relative to the hydrofluoroalkane
30 containing organic impurities. The amount of metal ion is more frequently at least 0,1 ppm. The amount is preferably at least 0,5 ppm.

The metal ion can be introduced into the reaction medium for example by addition of a suitable metal compound. In a particular embodiment, the treatment with chlorine is carried out in a reactor made of a material containing a suitable metal
35 as described above, under conditions sufficient to release at least a trace amount of metal ion.

In the third variant of the first aspect of the process according to the invention, the treatment with chlorine is generally carried out at a temperature of greater than or equal to 0°C. The temperature is often greater than or equal to 20°C. Preferably, the temperature is greater than or equal to about 40°C. In this variant, the treatment with chlorine is generally carried out at a temperature of less than or equal to 200°C. The temperature is often less than or equal to 150°C. Preferably, the temperature is less than or equal to about 100°C.

In the third variant of the first aspect of the process according to the invention, the duration of the treatment with chlorine is generally greater than or equal to 1 h. The duration of the treatment with chlorine is often greater than or equal to 3 h. In the third variant of the first aspect of the process according to the invention, the duration of the treatment with chlorine is generally less than or equal to 20 h. Preferably, the duration of the treatment with chlorine is less than or equal to about 10 h.

The suitable pressures in the third variant of the first aspect of the process according to the invention, are the same as in the second variant of the first aspect of the process according to the invention.

In the third variant of the first aspect of the process according to the invention, the hydrofluoroalkane is suitably selected from the group consisting of 1,1,1,2,3,3,3-heptafluoropropane, 1,1,1,3,3,3-hexafluoropropane, 1,1,1,3,3-pentafluoropropane and 1,1,1,3,3-pentafluorobutane. It is preferably selected from 1,1,1,3,3-pentafluoropropane and 1,1,1,3,3-pentafluorobutane. Most preferably, the hydrofluoroalkane is 1,1,1,3,3-pentafluorobutane.

In another embodiment, the third variant of the first aspect of the process according to the invention can also advantageously be used for the chlorination of bulk chloro(fluoro) olefins such as described above or of fractions comprising a high amount of such chloro(fluoro) olefins.

The first aspect of the process according to the invention may be carried out in batchwise, semi-continuous or continuous mode. A continuous mode is preferred.

In the first aspect of the process according to the invention, the chlorination reactor and the distillation apparatus are preferably made of corrosion-resistant materials such as, in particular, alloys of the type such as MONEL, INCONEL or HASTELLOY.

In the first aspect of the process according to the invention, care is advantageously taken to ensure that the oxygen content in the chlorine is less than

1000 ppm by volume and preferably that it does not exceed 50 ppm by volume. To do this, the hydrofluoroalkane containing olefinic impurities may first be deaerated by sparging with an inert gas, for example nitrogen.

In the first aspect of the process according to the invention, the treatment
5 with chlorine is generally followed by a separation operation whose function is to separate the impurities from the hydrofluoroalkane, after they have been chlorinated. The separation operation is preferably a distillation.

The first aspect of the process according to the invention applies in a most particularly preferred manner to the production of purified 1,1,1,3,3-
10 pentafluorobutane.

In a second aspect of the process according to the invention, the hydrofluoroalkane containing organic impurities is subjected to a reaction with hydrogen fluoride.

The second aspect makes it possible in particular to effectively reduce the
15 content of organic impurities present in the hydrofluoroalkane by using hydrogen fluoride. The latter compound is among reagents used in a synthesis of a hydrofluoroalkane by hydrofluorination. The products of the conversion are saturated (hydro)fluoroalkanes which are toxicologically and environmentally more acceptable than olefinic or chlorofluoro organic impurities. In addition, for
20 certain organic impurities, the reaction with hydrogen fluoride will lead to the formation of the desired hydrofluoroalkane, namely, in particular, 1,1,1,3,3-pentafluoropropane or 1,1,1,3,3-pentafluorobutane. The second aspect of the process according to the invention may be carried out readily by using technical means developed for reactions for the synthesis of hydrofluoroalkanes by
25 hydrofluorination.

The organic impurities whose content may be reduced in particular in the second aspect of the process according to the invention comprise at least one chlorine atom, such as chlorodifluoropropanes and chlorofluorobutanes or -
butenes. They are in particular (chloro)fluoro olefins containing 3 or 4 carbon
30 atoms, such as monochlorotrifluorobutene isomers.

The second aspect of the process according to the invention is also particularly useful for the elimination of the hydrofluoroalkenes mentioned above.

In the second aspect of the process according to the invention, the reaction of the hydrofluoroalkane with hydrogen fluoride is preferably carried out in the
35 presence of a fluorination catalyst. It may also be carried out in the absence of catalyst.

When the reaction of the hydrofluoroalkane with hydrogen fluoride is carried out in the presence of a catalyst, catalysts which can promote the addition of HF to an olefin and/or the replacement of a chlorine atom with a fluorine atom may be used. Among the catalysts which may be used, mention may be made of derivatives of metals chosen from the metals from groups IIIa, IVa and b, Va and b and VIb of the Periodic Table of the Elements (IUPAC, 1970) and mixtures thereof. Titanium, tantalum, molybdenum, boron, tin and antimony derivatives are more especially selected. Preferably, titanium or tin derivatives are used. Metal derivatives which may be mentioned are salts and more particularly halides. Preferably, the choice is made from chlorides, fluorides and chlorofluorides. Catalysts that are particularly preferred in the process for preparing the hydrofluoroalkane according to the invention are the chlorides, fluorides and chlorofluorides of titanium and of tin and mixtures thereof. Titanium tetrachloride and tin tetrachloride are particularly suitable for use.

In the second aspect of the process according to the invention, the molar ratio between the hydrogen fluoride and the organic impurities present in the hydrofluoroalkane is generally at least 1 mol/mol. Preferably, the process is performed with a molar ratio of at least 1.5 mol/mol. The molar ratio between the hydrogen fluoride and the organic compound used generally does not exceed 1000 mol/mol. It is preferable for this molar ratio not to exceed 10 mol/mol. In the second aspect of the process according to the invention, a molar ratio between the hydrogen fluoride and the olefinic impurities of not more than 3 is often maintained.

The second aspect of the process according to the invention may be carried out in batchwise or continuous mode.

When the second aspect of the process according to the invention is carried out in batchwise mode, the duration of the reaction of the hydrofluoroalkane containing organic impurities with hydrogen fluoride generally ranges from 10 min to 5 h. Preferably, this duration is at least 0.5 h. Advantageously, this duration is at least 1 h. In general, this duration does not exceed 4 h. Preferably, this duration does not exceed 2.5 h.

When the second aspect of the process according to the invention is carried out in continuous mode, the residence time of the reagents in the reactor is generally at least 0.5 h. Usually it does not exceed 30 h. Typically it ranges from 5 to 25 h. Preferably, it ranges from 10 to 20 h. The expression "residence time of the reagents in the reactor" is intended to denote the ratio between the volume of

the reaction medium and the flow rate by volume of the reaction medium at the reactor outlet.

In a first variant, which is preferred, the reaction of the hydrofluoroalkane containing organic impurities with hydrogen fluoride in the second aspect of the process according to the invention is carried out in the liquid phase. In this
5 variant, the temperature at which the reaction of the hydrofluoroalkane containing organic impurities with hydrogen fluoride is carried out is generally at least 60°C. Preferably, the temperature is at least 80°C. In general, the temperature does not exceed 160°C. Preferably, it does not exceed 140°C.

10 In this variant, the pressure is chosen so as to keep the reaction medium in liquid form. The pressure used varies as a function of the temperature of the reaction medium. It is generally less than or equal to 40 bar. Preferably, it is less than or equal to 35 bar. In a particularly advantageous manner, the pressure is less than or equal to 25 bar. In general, the pressure is greater than or equal to 5 bar.
15 Preferably, the pressure is greater than or equal to 10 bar.

In a second variant, the second aspect of the process according to the invention is carried out in the gas phase. This variant is particularly suitable for purifying 1,1,1,3,3-pentafluoropropane and 1,1,1,3,3-pentafluorobutane.

Specifically, 1,1,1,3,3-pentafluoropropane and 1,1,1,3,3-
20 pentafluorobutane, and in particular 1,1,1,3,3-pentafluorobutane, show surprising thermal stability, which allows them to be purified in the gas phase.

In this second variant, a fluorination catalyst based on a metal oxide chosen from chromium oxide, zirconium oxide and aluminium oxide, and mixtures thereof, is often used. Often, the metal oxide has a specific surface area
25 determined according to the BET method of at least 100 m²/g and preferably of at least 150 m²/g. Generally, this specific surface area is not more than 400 m²/g. The metal oxide is preferably amorphous.

In this second variant, the temperature of the reaction with hydrogen fluoride is generally at least 50°C. Preferably, the temperature is at least 100°C.
30 Generally, the temperature is not more than 400°C. Preferably, the temperature is not more than 300°C.

The second aspect of the process according to the invention finds an advantageous application to the purification of a hydrofluoroalkane obtained by synthesis by hydrofluorination, in particular by hydrofluorination of a
35 chloro(fluoro)carbon. In the latter case, it may be advantageous to reduce the

hydrogen chloride content of the hydrofluoroalkane containing organic impurities prior to its use in the second aspect of the process according to the invention.

The second aspect of the process according to the invention often comprises at least one subsequent treatment step intended to recover the hydrofluoroalkane. Examples of treatment steps which may be used are, inter alia, treatments which may be used to separate the residual hydrogen fluoride from the hydrofluoroalkane, such as, for example, adsorption onto a solid, for instance NaF or alumina, washing with water, an extraction operation, a separation by means of a suitable membrane, an extractive distillation or a distillation.

In a third aspect of the invention, the hydrofluoroalkane containing organic impurities is subjected to a distillation and the purified hydrofluoroalkane is removed from the top of the distillation column or from the side.

It has been found, surprisingly, that organic impurities present in the hydrofluoroalkane, in particular (hydro)(chloro)fluorocarbons comprising 3 or 4 carbon atoms, do not have a tendency to form an azeotrope with the hydrofluoroalkane and can thus be separated by means of the third aspect of the process according to the invention. The third aspect of the process according to the invention may be carried out easily.

The organic impurities whose content may be reduced by the third aspect of the process according to the invention generally comprise 3 or 4 carbon atoms. They are in particular hydrochlorofluoroalkanes and (chloro)fluoro olefins containing 3 or 4 carbon atoms, such as monochlorotrifluorobutene isomers. The third aspect of the process according to the invention is also particularly useful for the elimination of the hydrofluoroalkenes mentioned above.

The distillation pressure is generally less than 10 bar absolute. It is usually not more than 5 bar. Preferably, it is not more than 3 bar. Generally, the distillation pressure is at least 0.5 bar. It is usually at least 1 bar. Preferably, it is at least 1.5 bar.

In the present description of the third aspect of the process according to the invention, any reference to the pressure corresponds to the absolute pressure measured at the top of the distillation column.

The temperature at which the distillation is carried out corresponds approximately to the boiling point of the hydrofluoroalkane at the chosen pressure.

When the hydrofluoroalkane is 1,1,1,3,3-pentafluorobutane, good results are obtained at a pressure of about 1.5 to 3 bar and a temperature of about 50 to 70°C.

The distillation may be carried out in one or more distillation columns.
5 Preferably, only one column will be used.

The distillation columns which may be used are known per se. It is possible to use, for example, conventional plate columns or "dual-flow" plate columns or columns with bulk or structured packing.

The number of theoretical plates in the distillation is generally at least 10.
10 It is usually at least 15. A number of at least 20 gives good results.

The feed of hydrofluoroalkane containing organic impurities in the third aspect of the process according to the invention is generally carried out at a level below 50% of the number of theoretical plates of the column, it being understood that the top of the column corresponds to 100% of the number of theoretical
15 plates. This level is usually not more than 45% of the number of theoretical plates of the column. Generally, the feed is carried out at a level of at least 5% of the number of theoretical plates of the column. This level is usually at least 10% of the number of theoretical plates of the column.

If a side removal is carried out, it is generally carried out at the level
20 which corresponds to at least 50% of the number of theoretical plates of the distillation. The side removal is generally carried out at the level which corresponds to not more than 80% of the number of theoretical plates of the distillation.

In the third aspect of the process according to the invention, the purified
25 hydrofluoroalkane is generally removed in an amount of at least 50% of the feed. The amount is usually at least 70% of the feed. The amount is preferably at least 80% of the feed. Generally, the purified hydrofluoroalkane is removed in an amount of not more than 99% of the feed. The amount is usually not more than 97% of the feed. The amount is preferably not more than 95% of the feed.

30 The degree of molar reflux in the distillation is generally not more than 20. This degree is usually not more than 10. A degree of reflux of not more than 7 has given good results.

In a fourth aspect of the process according to the invention, the purification treatment is an extractive distillation. The extractive distillation is
35 carried out in the presence of at least one extractant which is generally chosen from (hydro)chlorocarbons, (hydro)fluorocarbons, hydrochlorofluorocarbons,

hydrocarbons, ketones, alcohols, ethers, esters, nitriles, hydrogen chloride and carbon dioxide.

Hydrofluorocarbons which may be used as extractants comprise typically from 1 to 6 carbon atoms, preferably from 1 to 4 carbon atoms. Preferred specific
5 hydrofluorocarbon extractants are hydrofluoroalkane extractants chosen, for example, from difluoromethane, 1,1,1,2-tetrafluoroethane, 1,1,1-trifluoroethane, pentafluoroethane, 1,1,1,2,3,3,3-heptafluoropropane 1,1,1,3,3-pentafluoropropane and 1,1,1,3,3-pentafluorobutane. It is understood that the hydrofluorocarbon extractant in the fourth aspect of the process according to the invention is, in
10 general, different from the hydrofluoroalkane containing organic impurities.

Other extractants which may be used are chosen, for example, from dichloromethane, perchloroethylene, n-pentane, n-hexane, methanol, ethanol, isopropanol, diethyl ether, acetone, 2-butanone, ethyl acetate and acetonitrile.

In another embodiment, the extractant is chosen from chlorinated
15 precursors suitable for a synthesis of the hydrofluoroalkane by hydrofluorination or from chloro(fluoro) intermediates obtainable by hydrofluorination of a said chlorinated precursor, such as chlorofluoropropanes and chlorofluorobutanes.

Preferably, the extractant is chosen from 1,1,1,3,3-pentachlorobutane, 1,1-dichloro-1,3,3-trifluorobutane, 1,3-dichloro-1,1,3-trifluorobutane, 3,3-dichloro-
20 1,1,1-trifluorobutane, 1-chloro-1,3,3,3-tetrafluorobutane and 3-chloro-1,1,3,3-tetrafluorobutane or a mixture of these extractants.

The distillation is generally carried out at a pressure and a temperature which makes it possible essentially to avoid, where appropriate, the formation of azeotropes between the extractant and the hydrofluoroalkane.

25 The distillation may be performed in one or more distillation columns. Preferably, only one column will be used.

The distillation columns which may be used in the fourth aspect of the process according to the invention are known per se. It is possible to use, for example, conventional plate columns or "dual-flow" plate columns or columns
30 with bulk or structured packing.

In a fifth aspect of the process according to the invention, the purification treatment is an adsorption onto a solid adsorbent. The solid adsorbent may be chosen, for example, from aluminas, silicas, iron oxide compounds, zeolites and active charcoals. Such adsorbents are commercially available. The adsorbent is
35 optionally activated prior to its use in the adsorption treatment. A heat treatment or a treatment intended to increase the Lewis acidity of the solid adsorbent is

suitable. The preferred solid adsorbents are those which have undergone a treatment intended to increase their Lewis acidity, for example a washing with hydrochloric acid or with nitric acid.

5 The contact between the hydrofluoroalkane containing organic impurities and the solid adsorbent may be carried out according to various techniques. The process may be performed in a fluidized bed, but it is generally preferred to place the solid adsorbent in the form of a fixed bed of particles, through which is passed a flow of the hydrofluoroalkane containing organic impurities. This flow may be liquid or gaseous. In one variant, the adsorption is carried out in the gas phase.

10 When the fifth aspect of the process is carried out in the gas phase, a contact time between the hydrofluoroalkane containing organic impurities and the solid adsorbent of at least 1 s is carried out. Preferably, the process is performed with a contact time of greater than 2 s. Good results have been obtained with a contact time of greater than or equal to 3 s. In principle, the process may be
15 performed with a very long contact time, for example of several minutes. In practice, for reasons of efficiency, the process is generally performed with a contact time of less than 1 minute and preferably less than or equal to about 30 s.

When the fifth aspect of the process is carried out in the liquid phase, a contact time between the hydrofluoroalkane containing organic impurities and the
20 solid adsorbent of at least about 2 minutes is carried out. Preferably, the process is performed with a contact time of greater than about 5 minutes.

In principle, the process may be performed with a very long contact time, for example of 120 minutes. In practice, the process is generally performed with a contact time of less than 60 minutes and preferably less than or equal to about 30
25 minutes.

When the fifth aspect of the process is carried out in a fixed bed, the contact time is defined as the ratio of the volume of the bed of adsorbent to the flow rate by volume of the stream of hydrofluoroalkane containing organic impurities. When the fifth aspect of the process is carried out in a fluidized bed,
30 the contact time is defined as the ratio of the volume of the tank containing the solid adsorbent to the flow rate by volume of the stream of hydrofluoroalkane containing organic impurities.

The solid adsorbent is used in the form of a powder of particles whose optimum particle size depends on the conditions under which the process is
35 carried out. In general, a solid adsorbent whose particle diameter ranges from about 0.1 mm to 10 mm is selected. The process is preferably performed with

particles with a diameter of less than or equal to 7 mm. In a particularly preferred manner, particles with a diameter of less than or equal to 5 mm are used.

Moreover, it is preferred to use a solid adsorbent whose particles have a diameter of greater than or equal to 0.5 mm. The process is preferably performed with
5 particles with a diameter of greater than or equal to 1 mm. In a particularly preferred manner, particles with a diameter of greater than or equal to 2 mm are used.

After the process, the solid adsorbent may be regenerated by heating at moderate temperature, for example 100 to 250°C, under a stream of gas, for
10 example under nitrogen, or under reduced pressure. The solid adsorbent may also be regenerated by a treatment with oxygen.

In a sixth aspect of the process according to the invention, the purification treatment is a reaction with a compound containing oxygen. It has been found that reagents containing oxygen react preferentially with the organic impurities present
15 in the hydrofluoroalkane, in particular in 1,1,1,3,3-pentafluorobutane and essentially without degrading the hydrofluoroalkane, in particular 1,1,1,3,3-pentafluorobutane. The compound containing oxygen may be, for example, an oxygenated gas, an oxygenated acid, an organic or inorganic peroxide, a peroxide salt or a peracid. Specific examples of such compounds are chosen from oxygen,
20 ozone, hydrogen peroxide, peracetic acid, potassium permanganate, sulphuric acid and sulphur trioxide.

In another embodiment of the sixth aspect of the process according to the invention, the reaction is carried out in the presence of a base and the compound containing oxygen is an alcohol. The base may be, for example, an alkali metal
25 hydroxide such as sodium hydroxide or potassium hydroxide. The alcohol may be chosen, for example, from methanol, ethanol and isopropanol.

The reaction with the compound containing oxygen may be carried out in the presence or in the absence of an oxygenation catalyst. Oxygenation catalysts which may be used may be chosen, for example, from compounds and in
30 particular from complexes containing platinum, manganese or titanium.

The reaction with the compound containing oxygen may be carried out in the gas phase or in the liquid phase. It is preferably carried out in the liquid phase. In this case, the reaction temperature is generally not more than 150°C. The temperature is more frequently not more than 120°C. Preferably, the temperature
35 is not more than 100°C. The reaction temperature is generally at least -20°C. The

temperature is more frequently at least 0°C. Preferably, the temperature is at least 20°C.

The reaction pressure is generally from 1 to 10 bar.

In a seventh aspect of the process according to the invention, the
5 purification treatment is a reaction in the gas phase with a reagent capable of reacting with at least some of the organic impurities, with the exception of a reaction with elemental chlorine.

In the seventh aspect of the process according to the invention, the reagent
may in principle be any reagent capable of reacting in the gas phase with at least
10 some of the organic impurities present in the hydrofluoroalkane and in particular with the olefinic impurities. The reagent is advantageously chosen from hydrogen chloride, hydrogen, hydrogen fluoride, oxygen and ozone.

In a typical example, the reaction is a catalytic hydrogenation.

It has been found, surprisingly, that catalytic hydrogenation makes it
15 possible to reduce the content of any impurity in particular in 1,1,1,3,3-pentafluorobutane to a level close to, even less than, 5 mg/kg, while at the same time avoiding degradation of the hydrofluoroalkane, in particular of 1,1,1,3,3-pentafluorobutane.

Catalysts which may be used in the catalytic hydrogenation reaction in the
20 gas phase according to the invention are, for example, catalysts containing a metal from group VIII of the Periodic Table of Elements (IUPAC, 1970) or a mixture of several metals, preferably supported on a support such as active charcoal, a fluorinated alumina or aluminium trifluoride. Specific examples of metals from group VIII are platinum, palladium and rhodium. Among these catalysts, a
25 catalyst comprising palladium is preferred.

The metal content in the supported catalysts which may be used is generally at least 0.001% by weight. This content is usually at least 0.1% by weight. The metal content in the supported catalysts is generally not more than 20% by weight. This content is frequently not more than 10% by weight. A
30 catalyst which is resistant with respect to the products which may be present during the catalytic hydrogenation, in particular hydrogen fluoride, is preferably chosen. Good results are obtained, for example, with a catalyst comprising palladium supported on active charcoal.

The molar ratio between the reagent and the organic impurities present in
35 the hydrofluoroalkane is generally at least 1 mol/mol. Preferably, the process is performed with a molar ratio of at least 1.5 mol/mol. The molar ratio between the

reagent and the organic impurities generally does not exceed 1000 mol/mol. It is preferable for this molar ratio not to exceed 10 mol/mol. In the seventh aspect of the process according to the invention, a molar ratio between the reagent and the olefinic impurities of not more than 3 is frequently maintained. However, when
5 the reagent is hydrogen, good results are also obtained when a molar ratio between the hydrogen and the olefinic impurities of greater than or equal to 5 is maintained. The molar ratio between the hydrogen and the olefinic impurities is advantageously less than or equal to 20. Preferably, this ratio is less than or equal to 10.

10 The temperature of the gas-phase reaction is generally at least 50°C. This temperature is usually at least 70°C. Preferably, this temperature is greater than or equal to 100°C. Generally, the temperature of the gas-phase reaction is not more than 400°C. Preferably, this temperature is not more than 300°C. In a particularly preferred manner, this temperature is not more than 250°C. Even more preferably,
15 this temperature is not more than 150°C.

In the seventh aspect of the process according to the invention, it is often necessary to carry out an operation intended to place the hydrofluoroalkane containing the organic impurities into the gaseous form. This operation may comprise, for example, an evaporation. In one preferred variant, the operation
20 comprises the removal, in the gaseous form, of a distillation fraction comprising hydrofluoroalkane and organic impurities, for the purpose of purifying it in the gas phase. The distillation fraction may be obtained by one or more distillations of crude hydrofluoroalkane comprising, in addition to organic impurities, possibly reagents arising as by-products or intermediates of the synthesis of the hydro-
25 fluoroalkane. The crude hydrofluoroalkane may in particular comprise hydrogen fluoride and/or hydrogen chloride, in particular when the hydrofluoroalkane is obtained by hydrofluorination. The hydrogen fluoride and/or hydrogen chloride content in the crude hydrofluoroalkane may be reduced by distillation, such that the distillation fraction has a low hydrogen fluoride and/or hydrogen chloride
30 content.

This reduction of the hydrogen fluoride and/or hydrogen chloride content is particularly advantageous when a catalytic hydrogenation as described above is carried out. In this case, hydrofluoroalkane containing organic impurities and having an acidity of not more than 1000 mmol/kg is generally used in the
35 purification treatment. Preferably, the acidity is not more than 100 mmol/kg.

Good results are obtained with hydrofluoroalkane containing organic impurities that are essentially free of hydrogen fluoride and/or hydrogen chloride.

In the operation intended to place the hydrofluoroalkane containing the organic impurities into the gaseous form, care is generally taken to ensure that the temperature of the hydrofluoroalkane does not exceed the temperature of the gas-phase purification treatment.

In the seventh aspect of the process according to the invention, the gas-phase purification reaction may be advantageously followed by one or more treatments intended to separate the hydrofluoroalkane from the products of reaction between the organic impurities and the reagent. A distillation is suitable as a treatment, in particular when the reagent is hydrogen.

In the process according to the invention, the purification treatment may be followed by one or more finishing steps intended, for example, to remove any residual acidity, in particular traces of hydrogen fluoride. A suitable finishing step for this purpose is, for example, an adsorption onto a solid such as alumina, NaF or silica.

Other treatments which may be used are, for example, a washing with water, an extraction operation or a separation by means of a suitable membrane.

The process according to the invention applies to the purification of a hydrofluoroalkane containing olefinic impurities, prepared by any synthetic process, without a pretreatment being required. The process according to the invention also applies to the purification of a hydrofluoroalkane containing organic impurities, which consists essentially of hydrofluoroalkane and organic impurities. Typically, the hydrofluoroalkane to be purified contains not more than 10% by weight of organic impurities. This content of impurities may be not more than 5% by weight. It may even be not more than 1% by weight. The process according to the invention may even be applied to a hydrofluoroalkane containing not more than 0.1% by weight of organic impurities.

The process according to the invention finds an advantageous application in the purification of a hydrofluoroalkane obtained by hydrofluorination, in particular by hydrofluorination of a hydrochloro(fluoro)carbon.

It should be understood that the different aspects of the invention can be combined with each other or with other purification treatments in order to optimise the benefits achieved by the process according to the invention. In a particular embodiment, the process according to the invention comprises 2, 3 or 4 purification steps for removing organic impurities, including at least one

purification treatment according to the invention. In particular the combinations allow for effective reduction of chloro(fluoro) olefin content with very low losses of desired hydrofluoroalkane.

In the following paragraph describing combinations of purification treatments, the following abbreviations are used:

- (a1) a treatment with chlorine according to the first variant of the first aspect of the process according to the invention;
- (a2) a treatment with chlorine according to the second variant of the first aspect of the process according to the invention;
- 10 (a3) a treatment with chlorine according to the third variant of the first aspect of the process according to the invention;
- (b) a reaction with hydrogen fluoride according to the second aspect of the process according to the invention;
- (c) a distillation according to the third aspect of the process according to the invention;
- 15 (d) an extractive distillation according to the fourth aspect of the process according to the invention;
- (e) an adsorption onto a solid adsorbent according to the fifth aspect of the process according to the invention;
- 20 (f) a reaction with a compound containing oxygen according to the sixth aspect of the process according to the invention;
- (g) a gas-phase reaction according to the seventh aspect of the process according to the invention;
- (h) a photochlorination using exclusively UV light of a wavelength > 280 nm
- 25 (i) a photolysis in the absence of chlorine
- (j) a reaction with fluorine

Suitable consecutive combinations include, amongst others, ("+" meaning "followed by")

- (a3) +(a1), (a3) +(a2), (a3) +(c), (a3) +(e), (a3) +(h), (b) +(a1), (b) +(a2), (b) +(c), (b) +(e), (b) +(h), (c) +(a1), (c) +(a2), (c) +(c), (c) +(e), (c) +(h), (d) +(a1), (d) +(a2), (d) +(c), (d) +(e), (d) +(h), (f) +(a1), (f) +(a2), (f) +(c), (f) +(e), (f) +(h), (g) +(a1), (g) +(a2), (g) +(c), (g) +(e), (g) +(h), (i) +(a1), (i) +(a2), (i) +(c), (i) +(e), (j) +(a1), (j) +(a2), (j) +(c), (j) +(e), (i) +(h) +(c).

- Combinations (a3) +(a1), (a3) +(a2), (a3) +(c), (a3) +(e), (a3) +(h), (c) +(a1), (c) +(a2), (c) +(c), (c) +(e), (c) +(h) (i) +(a1), (i) +(a2), (i) +(c), (i) +(e), (i) +(h) +(c) are preferred.

Combinations (a3) +(a1), (a3) + (a2), (a3) + (c), (a3) + (h), (c) +(a1), (c) + (a2), (c) + (c), (c) + (h), (i) +(h) +(c) are particularly preferred.

A variant of particularly preferred combinations includes (a3) +(a1) + (c), (a3) + (a2) + (c), (a3) + (c) + (c), (a3) + (h) + (c).

5 It is understood that the aforementioned combinations are particularly well suited for the purification of 1,1,1,3,3-pentafluorobutane.

The invention also relates to a purified hydrofluoroalkane, preferably chosen from 1,1,1,3,3-pentafluoropropane and 1,1,1,3,3-pentafluorobutane, which has a hydrofluoroalkene content of less than 30 ppm by weight and preferably less than
10 20 ppm. A hydrofluoroalkene content of less than 10 ppm is particularly preferred. When the hydrofluoroalkane is 1,1,1,3,3-pentafluorobutane, they are usually hydrofluoroalkenes of empirical formula $C_4F_4H_4$, mentioned above.

The examples which follow are intended to illustrate the present invention without, however, limiting its scope.

15 Example 1 – Chlorination initiated by electromagnetic radiation,
comprising at least one fraction of wavelengths less than 280 nm

750 g of 1,1,1,3,3-pentafluorobutane (HFC-365mfc) containing 709 ppm by weight of olefinic impurities including 40 ppm of $C_4H_4F_4$ hydrofluoroalkenes and chlorine were introduced into a tank with a working volume of 0.7 l equipped
20 with an immersion UV emitter of the type Heraeus TQ 150, the power of which was 150 W and the arc length of which was 44 mm. This low-pressure mercury vapour emitter has broadband emission in the range of UV rays above 190 nm. All the internal glassware between the emitter and the 1,1,1,3,3-pentafluorobutane was made of quartz. The molar ratio between the chlorine and the olefinic
25 impurities was 1.6. The lamp was cooled so as to maintain the 1,1,1,3,3-pentafluorobutane at a temperature of 25°C. The medium was stirred by means of a magnetic bar. Samples were taken at regular intervals for analysis of the olefinic impurity content of the 1,1,1,3,3-pentafluorobutane by gas chromatography.

The table below shows the change in concentration of the olefins in the
30 1,1,1,3,3-pentafluorobutane which was observed after a given period of treatment with chlorine in the presence of UV light comprising a fraction of wavelengths less than 280 nm.

| | 30 min | 1 h | 2 h |
|---|--------|-----|-----|
| C ₄ H ₄ F ₄ content (ppm/weight) | 9 | 5 | 0 |
| Total content of olefinic impurities (ppm/weight) | 83 | 18 | 0 |

Example 2 – Hydrofluorination

A reactor containing hydrogen fluoride and titanium tetrachloride was fed continuously with a 1,1,1,3,3-pentafluorobutane containing 5.8% by weight of two trifluorodichlorobutene isomers. The reactor was maintained at a temperature of about 130°C with continuous stirring. The pressure was adjusted to 22 bar. The purified 1,1,1,3,3-pentafluorobutane was removed continuously in the gas phase. The content of the first trifluorodichlorobutene isomer in the purified 1,1,1,3,3-pentafluorobutane was 0.1% by weight. The content of the second trifluorodichlorobutene isomer in the purified 1,1,1,3,3-pentafluorobutane was 0.05% by weight.

Example 3 – Distillation with removal from the top of the column

Distillation was carried out in a column packed with 5/8" "pall" rings. 1,1,1,3,3-Pentafluorobutane containing 94 ppm of 3 saturated chlorofluoro organic impurities containing 3 or 4 carbon atoms and 252 ppm of 2 (chloro)fluoro olefins containing 4 carbon atoms was fed in at the level of the column which corresponds to 43% of the number of theoretical plates. The pressure was 2 bar, the temperature was 60°C and the degree of molar reflux was about 4. An amount of purified 1,1,1,3,3-pentafluorobutane corresponding to 84% of the feed was removed from the top of the column. This product still contained 6 ppm of a chlorofluoro saturated organic impurity containing 3 carbon atoms. The content of the other two saturated impurities was reduced below the 5 ppm detection limit. The content of the said 2 (chloro)fluoro olefins containing 4 carbon atoms in the purified 1,1,1,3,3-pentafluorobutane was reduced to below the 5 ppm detection limit.

Example 4 – (comparative)

The distillation was carried out in the same apparatus as in Example 3. 1,1,1,3,3-Pentafluorobutane containing 170 ppm of 3 chlorofluoro saturated organic impurities containing 3 or 4 carbon atoms and 152 ppm of 2 chlorofluoro olefins containing 3 or 4 carbon atoms was fed into the level of the column which corresponds to 65% of the number of theoretical plates. The pressure was 2 bar,

the temperature was 60°C and the degree of molar reflux was about 25. An amount of 1,1,1,3,3-pentafluorobutane corresponding to 85% of the feed was removed from the bottom of the column. This product contained 170 ppm of the said 3 saturated organic impurities and 141 ppm of the said 2 chlorofluoro olefins containing 4 carbon atoms.

It appears that the third aspect of the process according to the invention allows a fast, efficient and complete purification of olefinic impurities, in particular of hydrofluoroalkenes, from the hydrofluoroalkane. No degradation of the hydrofluoroalkane HFC-365mfc was observed.

10 Example 5 – Reagent containing oxygen

1,1,1,3,3-Pentafluorobutane containing 3260 mg/kg of (chloro)fluoro olefinic impurities was mixed in the liquid phase with 96% sulphuric acid in a 1:1 1,1,1,3,3-pentafluorobutane/sulphuric acid ratio by volume. The mixture was stirred for 24 h at 40°C. After this period, the recovered 1,1,1,3,3-pentafluorobutane contained 140 ppm of (chloro)fluoro olefinic impurities.

15 Example 6 - Gas-phase reaction

1,1,1,3,3-Pentafluorobutane with a purity of 99.86% by weight, containing 755 mg/kg of (chloro)fluoro olefinic impurities, was evaporated at 85°C and fed continuously into a reactor, containing a catalyst consisting of 0.5% palladium on active charcoal. The reactor was also fed continuously with hydrogen. The molar ratio between the hydrogen and the 1,1,1,3,3-pentafluorobutane was 5.8×10^{-3} . The reaction temperature was 100°C. The gaseous effluents of the reaction were condensed with a water condenser at 5°C and were analysed by GC. They consisted of 1,1,1,3,3-pentafluorobutane with a purity of 99.92% by weight, containing less than 5 mg/kg of (chloro)fluoro olefinic impurities. No degradation of the 1,1,1,3,3-pentafluorobutane was observed.

CLAIMS

1 - Process for obtaining a hydrofluoroalkane comprising at least three carbon atoms, which is purified of organic impurities including (chloro)fluoro olefins, according to which the hydrofluoroalkane containing organic impurities is
5 subjected to at least one purification treatment chosen from

- (a) a treatment with chlorine in the presence of a initiator
- (b) a reaction with hydrogen fluoride
- (c) a distillation in which the purified hydrofluoroalkane is removed
from the top of the distillation column or from the side
- 10 (d) an extractive distillation
- (e) an adsorption onto a solid adsorbent
- (f) a reaction with a compound containing oxygen, and
- (g) a gas-phase reaction with a reagent capable of reacting with at least
some of the organic impurities, with the exception of a reaction with elemental
15 chlorine.

2 - Process according to Claim 1, in which the hydrofluoroalkane containing the olefinic impurities is subjected to a treatment with chlorine in the presence of an initiator.

3 - Process according to Claim 2, in which the hydrofluoroalkane is
20 chosen from 1,1,1,3,3-pentafluoropropane and 1,1,1,3,3-pentafluorobutane.

4 - Process according to Claim 2 or 3, in which the initiator is an organic initiator preferably chosen from peroxide and diazo compounds.

5 - Process according to Claim 4, characterized in that the organic initiator is chosen from peroxide compounds.

25 6 - Process according to Claim 5, characterized in that the peroxide compound is chosen from diacyl peroxides and peroxydicarbonates.

7 - Process according to any one of Claims 2 to 6, characterized in that the initiator is selected from compounds with a half-life of about 1 hour at the temperature of the treatment with chlorine.

8 - Process according to any one of Claims 2 to 7, characterized in that the treatment with chlorine is carried out in the presence of at least about 10 ppm by weight of organic initiator relative to the weight of hydrofluoroalkane containing the olefinic impurities.

5 9 - Process according to Claim 2 or 3, in which the initiator is an electromagnetic radiation comprising at least one fraction of wavelengths less than 280 nm.

10 10 - Process according to Claim 9, in which the energy of the fraction of wavelengths less than 280 nm is at least 10% of the total energy of the electromagnetic radiation.

11 - Process according to Claim 10, in which the energy of the fraction of wavelengths less than 280 nm is at least 20% of the total energy of the electromagnetic radiation.

15 12 - Process according to any one of Claims 9 to 11, in which the wavelengths in the fraction of wavelengths less than 280 nm are essentially at least 170 nm.

13 - Process according to any one of Claims 9 to 12, in which the molar ratio between the chlorine and the sum of the olefinic impurities present is from 1 to 10.

20 14 - Process according to any one of Claims 9 to 12, in which the molar ratio between the chlorine and the sum of the olefinic impurities present is less than 1.

25 15 - Process according to claim 2 or 3, in which the initiator is a low amount of metal ion, preferably selected from ions of group IIIa, IVa and b, Va and b, VIB and VIII metals.

16 - Process according to any one of Claims 2 to 15, characterized in that the treatment with chlorine is carried out in the liquid phase.

17 - Process according to Claim 1, in which the hydrofluoroalkane containing organic impurities is subjected to a reaction with hydrogen fluoride.

18 - Process according to Claim 17, in which the reaction with hydrogen fluoride is carried out in the presence of a fluorination catalyst.

19 - Process according to Claim 17 or 18, in which the reaction with hydrogen fluoride is carried out in the liquid phase.

5 20 - Process according to Claim 18 or 19, in which the fluorination catalyst is chosen from compounds of metals from groups IIIa, IVa, IVb, Va, Vb and VIb.

21 - Process according to Claim 17 or 18, in which the reaction with hydrogen fluoride is carried out in the gas phase.

10 22 - Process according to Claim 21, in which the fluorination catalyst is based on a metal oxide chosen from chromium oxide, zirconium oxide and aluminium oxide, and mixtures thereof.

23 - Process according to Claim 21 or 22, in which the temperature of the reaction with hydrogen fluoride is from 50°C to 400°C.

15 24 - Process according to any one of Claims 17 to 23, in which a molar ratio between the hydrogen fluoride and the olefinic impurities of not more than 3 is maintained.

25 25 - Process according to Claim 1, in which the hydrofluoroalkane containing organic impurities is subjected to a distillation and the purified hydrofluoroalkane is removed from the top of the distillation column or from the side.

26 - Process according to Claim 25, in which the purified hydrofluoroalkane is removed from the top of the distillation column.

25 27 - Process according to Claim 25, in which the purified hydrofluoroalkane is removed from the side.

28 - Process according to any one of Claims 25 to 27, in which the distillation pressure is from 0.5 to 10 bar.

29 - Process according to any one of Claims 25 to 28, in which the temperature at which the distillation is carried out corresponds approximately to the boiling point of the hydrofluoroalkane at the chosen pressure.

30 - Process according to any one of Claims 25 to 29, in which the degree
5 of molar reflux of the distillation is not more than 20.

31 - Process according to any one of Claims 25 to 30, in which the feed of hydrofluoroalkane containing organic impurities is carried out at a level of from 5% to 50% of the number of theoretical plates of the column.

32 - Process according to any one of Claims 25 and 27 to 31, in which the
10 side removal is carried out at the level which corresponds to from 50% to 80% of the total number of theoretical plates of the distillation.

33 - Process according to Claim 1, in which the purification treatment is an extractive distillation.

34 - Process according to Claim 33, in which the extractive distillation is
15 carried out in the presence of at least one extractant chosen from (hydro)chlorocarbons, (hydro)fluorocarbons, hydrochlorofluorocarbons, hydrocarbons, ketones, alcohols, ethers, esters, nitriles, hydrogen chloride and carbon dioxide.

35 - Process according to Claim 34, in which the extractant is chosen from 1,1,1,3,3-pentachlorobutane, 1,1-dichloro-1,3,3-trifluorobutane, 1,3-dichloro-
20 1,1,3-trifluorobutane, 3,3-dichloro-1,1,1-trifluorobutane, 1-chloro-1,3,3,3-tetrafluorobutane and 3-chloro-1,1,3,3-tetrafluorobutane or a mixture of these extractants.

36 - Process according to Claim 1, in which the purification treatment is an adsorption onto a solid adsorbent.

37 - Process according to Claim 36, in which the solid adsorbent is chosen
25 from aluminas, iron oxides, silicas, zeolites and active charcoals, that are optionally activated.

38 - Process according to Claim 36 or 37, in which the adsorption is carried out in the gas phase.

39 - Process according to Claim 1, in which the purification treatment is a reaction with a compound containing oxygen.

40 - Process according to Claim 39, in which the compound containing oxygen is chosen from oxygen, ozone, hydrogen peroxide, peracetic acid,
5 potassium permanganate, sulphuric acid and sulphur trioxide.

41 - Process according to Claim 39, in which the reaction is carried out in the presence of a base and the compound containing oxygen is an alcohol.

42 - Process according to any one of Claims 39 to 41, in which the reaction is carried out in the liquid phase.

10 43 - Process according to Claim 1, in which the purification treatment is a reaction in the gas phase with a reagent capable of reacting with at least some of the organic impurities, with the exception of a reaction of elemental chlorine with unsaturated impurities under irradiation with UV light with a wavelength of at least 280 nm.

15 44 - Process according to Claim 43, in which the reagent is chosen from chlorine, hydrogen chloride, hydrogen, hydrogen fluoride, oxygen and ozone.

45 - Process according to Claim 44, in which the reaction is a catalytic hydrogenation.

20 46 - Process according to any one of Claims 43 to 45, in which the reaction temperature is not more than 300°C.

47 - Process according to any one of Claims 43 to 46, comprising an operation of placing the 1,1,1,3,3-pentafluorobutane containing organic impurities into the gaseous form, the said operation being carried out at a temperature which does not exceed the temperature of the gas-phase purification treatment.

25 48 - Process according to claim 1 which comprises 2, 3 or 4 purification steps for removing organic impurities, including at least one purification treatment according to any one of claims 2 to 47.

49 - Process according to any one of Claims 1 to 48, in which the olefinic impurities comprise chlorofluoro olefins containing 3 or 4 carbon atoms.

50 - Process according to any one of Claims 1 to 49, in which the olefinic impurities comprise at least one hydrofluoroalkene.

51 - Process according to any one of Claims 1 to 50, in which the hydrofluoroalkane is chosen from 1,1,1,3,3,3-hexafluoropropane (HFC-236fa),
5 1,1,1,2,3,3,3-heptafluoropropane (HFC-227ea) and 1,1,1,2,3,4,4,5,5,5-decafluoropentane (HFC-43-10mee).

52 - Process according to any one of Claims 1 to 51, in which the hydrofluoroalkane is 1,1,1,3,3-pentafluorobutane.

53 - Purified hydrofluoroalkane chosen from 1,1,1,3,3-pentafluoropropane
10 and 1,1,1,3,3-pentafluoro-butane, which has a hydrofluoroalkene content of less than 30 ppm by weight.

54 - Purified hydrofluoroalkane according to Claim 53, with a hydrofluoroalkene content of less than 20 ppm.

INTERNATIONAL SEARCH REPORT

In **International Application No**
PCT/EP 01/10064

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07C17/395 C07C17/383 C07C17/386 C07C17/389 C07C19/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|---|---|
| X | US 5 944 962 A (BOYCE C BRADFORD) 31 August 1999 (1999-08-31) the whole document --- | 1-3,9, 16,48, 49,51,52 |
| X | WO 97 37955 A (ALLIED SIGNAL INC) 16 October 1997 (1997-10-16) the whole document --- | 1-3,9, 13,16,49 |
| X | DE 199 40 104 A (SOLVAY FLUOR & DERIVATE) 9 March 2000 (2000-03-09) cited in the application A the whole document --- | 1,36,37, 43-46, 49,50, 52-54 1,2,17 |
| X | EP 0 511 612 A (HOECHST AG) 4 November 1992 (1992-11-04) claims --- | 1,36-38, 49-51 |
| | -/-- | |

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

8 January 2002

Date of mailing of the international search report

16/01/2002

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

Bonnevalle, E

INTERNATIONAL SEARCH REPORT

In International Application No.

PCT/JP 01/10064

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|---|-------------------------|
| X | EP 0 733 612 A (HOECHST AG) 25 September 1996 (1996-09-25) the whole document | 1, 36-38, 50, 51 |
| X | ----- CHEMICAL ABSTRACTS, vol. 118, no. 19, 10 May 1993 (1993-05-10) Columbus, Ohio, US; abstract no. 191166, YOSHINAGA, MASAMI ET AL: "Purification of hydrochlorofluorocarbons and hydrofluorocarbons" XP000354050 abstract & JP 04 300842 A (SHOWA DENKO K. K., JAPAN) 23 October 1992 (1992-10-23) | 1, 39, 40, 48, 50 |
| X | ----- EP 0 512 502 A (HOECHST AG) 11 November 1992 (1992-11-11) the whole document | 1, 39, 41, 42, 48-51 |
| X | ----- EP 0 811 591 A (ATOCHM ELF SA) 10 December 1997 (1997-12-10) column 3, line 27 - line 30; claim 1 | 1, 39, 40, 42 |
| X | ----- EP 0 574 756 A (HOECHST AG) 22 December 1993 (1993-12-22) claims | 1, 39, 41, 42 |
| X | ----- EP 0 370 688 A (DU PONT) 30 May 1990 (1990-05-30) page 4, line 49 - page 5, line 9; claim 1 | 1, 39, 49, 50 |
| X | ----- WO 90 08750 A (DU PONT) 9 August 1990 (1990-08-09) claims | 1, 43-46, 50 |
| A | ----- EP 0 582 494 A (ATOCHM ELF SA) 9 February 1994 (1994-02-09) claims | 1, 17, 18, 21-23 |
| A | ----- US 5 895 825 A (CHEN BIN ET AL) 20 April 1999 (1999-04-20) column 2, line 29 - line 40; claims 1, 4 | 1, 25, 49 |
| A | ----- WO 00 29361 A (DAIKIN IND LTD) 25 May 2000 (2000-05-25) abstract -& EP 1 132 365 A (DAIKIN IND LTD) 12 September 2001 (2001-09-12) claim 3 | 1, 25, 49 |
| A | ----- US 4 950 364 A (WISMER JOHN A) 21 August 1990 (1990-08-21) claim 1 | 1, 33, 34 |
| | ----- -/-- | |

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 01/10064

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|--|-----------------------|
| A | US 3 101 304 A (WIIST HERBERT A) 20 August 1963 (1963-08-20) claims --- | 1, 33, 34 |
| A | WO 98 00381 A (ALLIED SIGNAL INC) 8 January 1998 (1998-01-08) claim 14 --- | 1, 33 |
| A | US 2 999 885 A (HEBERLING JR JACK W) 12 September 1961 (1961-09-12) claim 1 --- | 1, 39, 40 |
| P, X | DE 100 29 283 A (SOLVAY FLUOR & DERIVATE) 21 December 2000 (2000-12-21) claims; example 12 --- | 1-3, 9, 16, 49, 52 |
| E | WO 01 83411 A (INEOS FLUOR HOLDINGS LTD) 8 November 2001 (2001-11-08) the whole document ----- | 1, 36-38, 49-51 |

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

P/ 01/10064

| Patent document cited in search report | | Publication date | Patent family member(s) | Publication date |
|---|---|---------------------|----------------------------|---------------------|
| US 5944962 | A | 31-08-1999 | US 5750010 A | 12-05-1998 |
| WO 9737955 | A | 16-10-1997 | EP 0902776 A1 | 24-03-1999 |
| | | | JP 2000508315 T | 04-07-2000 |
| | | | WO 9737955 A1 | 16-10-1997 |
| | | | US 6077982 A | 20-06-2000 |
| DE 19940104 | A | 09-03-2000 | DE 19940104 A1 | 09-03-2000 |
| | | | AU 1147700 A | 27-03-2000 |
| | | | BR 9913453 A | 24-07-2001 |
| | | | CN 1316983 T | 10-10-2001 |
| | | | WO 0014040 A1 | 16-03-2000 |
| | | | EP 1109766 A1 | 27-06-2001 |
| EP 0511612 | A | 04-11-1992 | CA 2067078 A1 | 28-10-1992 |
| | | | DE 59207602 D1 | 16-01-1997 |
| | | | EP 0511612 A2 | 04-11-1992 |
| | | | ES 2097829 T3 | 16-04-1997 |
| | | | GR 3022734 T3 | 30-06-1997 |
| | | | JP 6056713 A | 01-03-1994 |
| | | | SG 43952 A1 | 14-11-1997 |
| | | | US 5233107 A | 03-08-1993 |
| EP 0733612 | A | 25-09-1996 | DE 19510159 A1 | 26-09-1996 |
| | | | AT 184269 T | 15-09-1999 |
| | | | DE 59602982 D1 | 14-10-1999 |
| | | | EP 0733612 A1 | 25-09-1996 |
| | | | ES 2139267 T3 | 01-02-2000 |
| | | | JP 8268932 A | 15-10-1996 |
| | | | US 5621152 A | 15-04-1997 |
| JP 4300842 | A | 23-10-1992 | NONE | |
| EP 0512502 | A | 11-11-1992 | DE 4115025 A1 | 12-11-1992 |
| | | | AT 150744 T | 15-04-1997 |
| | | | CA 2068128 A1 | 09-11-1992 |
| | | | DE 59208257 D1 | 30-04-1997 |
| | | | DK 512502 T3 | 22-09-1997 |
| | | | EP 0512502 A2 | 11-11-1992 |
| | | | ES 2101768 T3 | 16-07-1997 |
| | | | GR 3023875 T3 | 30-09-1997 |
| | | | JP 3133821 B2 | 13-02-2001 |
| | | | JP 5124999 A | 21-05-1993 |
| | | | SG 46681 A1 | 20-02-1998 |
| | | | US 5475169 A | 12-12-1995 |
| EP 0811591 | A | 10-12-1997 | FR 2749581 A1 | 12-12-1997 |
| | | | AU 712937 B2 | 18-11-1999 |
| | | | AU 2471597 A | 11-12-1997 |
| | | | CA 2207993 A1 | 06-12-1997 |
| | | | CN 1168875 A | 31-12-1997 |
| | | | DE 69704393 D1 | 03-05-2001 |
| | | | DE 69704393 T2 | 23-08-2001 |
| | | | EP 0811591 A1 | 10-12-1997 |
| | | | ES 2155658 T3 | 16-05-2001 |
| | | | JP 10059879 A | 03-03-1998 |
| | | | US 5801294 A | 01-09-1998 |

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/JP 01/10064

| Patent document cited in search report | | Publication date | Patent family member(s) | Publication date |
|---|---|---------------------|--|--|
| EP 0574756 | A | 22-12-1993 | AT 144971 T CA 2098245 A1 DE 59304392 D1 DK 574756 T3 EP 0574756 A1 ES 2096141 T3 JP 6065118 A SG 43281 A1 | 15-11-1996 14-12-1993 12-12-1996 14-04-1997 22-12-1993 01-03-1997 08-03-1994 17-10-1997 |
| EP 0370688 | A | 30-05-1990 | AT 108429 T AU 626349 B2 AU 4540289 A BR 8905864 A CA 2003039 A1 CN 1043118 A DE 68916756 D1 EP 0370688 A1 ES 2056229 T3 IN 171615 A1 JP 2223534 A MX 170445 B SU 1836312 A3 US 5449845 A US 5449846 A ZA 8908910 A | 15-07-1994 30-07-1992 31-05-1990 19-06-1990 22-05-1990 20-06-1990 18-08-1994 30-05-1990 01-10-1994 28-11-1992 05-09-1990 23-08-1993 23-08-1993 12-09-1995 12-09-1995 31-07-1991 |
| WO 9008750 | A | 09-08-1990 | US 5001287 A AT 107620 T AU 624480 B2 AU 5022490 A BR 9007057 A CA 2009064 A1 CN 1044647 A , B DE 69010192 D1 DE 69010192 T2 EP 0456691 A1 JP 4503064 T MX 166048 B WO 9008750 A1 ZA 9000803 A | 19-03-1991 15-07-1994 11-06-1992 24-08-1990 12-11-1991 02-08-1990 15-08-1990 28-07-1994 05-01-1995 21-11-1991 04-06-1992 16-12-1992 09-08-1990 30-10-1991 |
| EP 0582494 | A | 09-02-1994 | FR 2694556 A1 AU 666958 B2 AU 4445893 A CA 2100594 A1 CN 1085537 A , B DE 69300901 D1 DE 69300901 T2 EP 0582494 A1 ES 2080595 T3 GR 3018814 T3 JP 2509142 B2 JP 6184015 A KR 9604870 B1 | 11-02-1994 29-02-1996 10-02-1994 06-02-1994 20-04-1994 11-01-1996 01-08-1996 09-02-1994 01-02-1996 30-04-1996 19-06-1996 05-07-1994 16-04-1996 |
| US 5895825 | A | 20-04-1999 | DE 69801973 D1 EP 0919529 A1 JP 11228461 A | 15-11-2001 02-06-1999 24-08-1999 |

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PC P 01/10064

| Patent document cited in search report | | Publication date | | Patent family member(s) | Publication date |
|---|---|---------------------|------|----------------------------|---------------------|
| WO 0029361 | A | 25-05-2000 | EP | 1132365 A1 | 12-09-2001 |
| | | | WO | 0029361 A1 | 25-05-2000 |
| US 4950364 | A | 21-08-1990 | CA | 2000136 A1 | 04-11-1990 |
| | | | DE | 68912114 D1 | 17-02-1994 |
| | | | EP | 0395793 A1 | 07-11-1990 |
| | | | JP | 2295937 A | 06-12-1990 |
| | | | JP | 2619721 B2 | 11-06-1997 |
| US 3101304 | A | 20-08-1963 | DE | 1161877 B | |
| | | | FR | 1293690 A | 18-05-1962 |
| | | | GB | 910493 A | 14-11-1962 |
| | | | NL | 264123 A | |
| WO 9800381 | A | 08-01-1998 | US | 6001796 A | 14-12-1999 |
| | | | WO | 9800381 A1 | 08-01-1998 |
| | | | US | 6060629 A | 09-05-2000 |
| US 2999885 | A | 12-09-1961 | NONE | | |
| DE 10029283 | A | 21-12-2000 | DE | 10029283 A1 | 21-12-2000 |
| | | | AU | 6148300 A | 02-01-2001 |
| | | | WO | 0076945 A2 | 21-12-2000 |
| WO 0183411 | A | 08-11-2001 | WO | 0183411 A1 | 08-11-2001 |